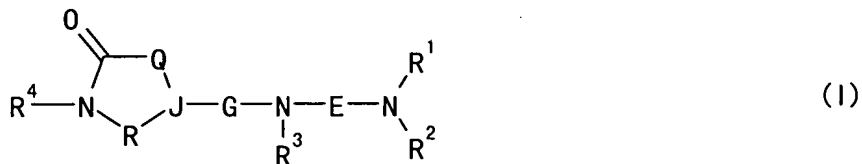


[illegible]

5 wherein R¹ is a hydrocarbon group, R² is a hydrocarbon group having 2 or more carbon atoms, where R¹ and R² may in combination form, together with an adjacent nitrogen atom, a ring optionally having a substituent or substituents, R³ is a hydrocarbon group optionally having a substituent or
10 substituents or a heterocyclic group optionally having a substituent or substituents, R⁴ is a hydrogen atom, a hydrocarbon group, a heterocyclic group and the like, E is a divalent chain hydrocarbon group and the like, G is CO or SO₂, J is a nitrogen atom, a methine group and the like, and
15 Q and R are each a divalent chain C₁₋₃ hydrocarbon group and the like, and a salt thereof show a superior CCR5 antagonistic activity and are useful as agents for the prophylaxis or treatment of HIV infection of human peripheral blood mononuclear cells, particularly AIDS.